TRANSMITTAL FORM  (to be used for all correspondence after initial for Total Number of Pages in This Submission	ro persons are required to respond to a colle	Approved for use through 07/31/2006. OMB 0651-0031 stent and Trademark Office; U.S. DEPARTMENT OF COMMERCE ction of information unless it displays a valid OMB control number.  10/661,966  September 11, 2003  Richard B. Roth  1645  TBA  SEQ-4038-UT
Fee Transmittal Form Fee Attached  Amendment/Reply After Final Affidavits/declaration(s) Extension of Time Request Express Abandonment Request Information Disclosure Statement  Certified Copy of Priority Document(s)	Drawing(s)  Licensing-related Papers  Petition Petition to Convert to a Provisional Application Power of Attorney, Revocation Change of Correspondence Actorney for Refund  Request for Refund  CD, Number of CD(s)  Landscape Table on CD  Remarks	After Allowance Communication to TC  Appeal Communication to Board of Appeals and Interferences  Appeal Communication to TC (Appeal Notice, Brief, Reply Brief)  Proprietary Information
Reply to Missing Parts/ Incomplete Application Reply to Missing Parts under 37 CFR 1.52 or 1.53		eg. No. 47,608

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This collection of information is required by 37 CFR 1.5. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.11 and1.14. This collection is estimated to 2 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.



### IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant:

ROTH, et al.

Examiner: Unknown

Serial No.:

10/661,966

Group Art Unit: 1645

Filed:

September 11, 2003

Docket: SEQ-4038-UT

Title:

METHODS FOR IDENTIFYING SUBJECTS AT RISK OF MELANOMA

AND TREATMENTS THEREOF

#### **COMMUNICATION RE: MISSING PARTS**

Mail Stop Missing Parts Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Dear Sir:

In response to the "Notice to File Missing Parts" please find computer-readable form of the Sequence Listing.

Applicants assume the application is now in proper order and in condition for examination. Please direct any inquiries to the undersigned attorney at 858 623-9470.

If there are any additional fees due or overpayment please contact the undersigned attorney at 858-623-9470.

Respectfully submitted,

Date: August 22, 2005 BioTechnology Law Group 658 Marsolan Avenue Solana Beach, CA 92075

Bruce D. Grant Reg. No. 47,608

CERTIFICATE OF MAILING PURSUANT TO 37 CFR § 1.8:

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APPLICATION NUMBER

FILING OR 371 (c) DATE

FIRST NAMED APPLICANT

ATTORNEY DOCKET NUMBER

10/661,966

MINNEAPOLIS, MN 55402

c/o PORTFOLIO IP

P.O. BOX 52050

**BIOTECHNOLOGY LAW GROUP** 

09/11/2003

Richard B. Roth

SEQ 4038 UT

**CONFIRMATION NO. 9006** 

FORMALITIES LETTER

\*OC000000016568254\*

Date Mailed: 07/20/2005

## NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

Filing Date Granted

Applicant is given **TWO MONTHS FROM THE DATE OF THIS NOTICE** within which to file the items indicated below to avoid abandonment. Extensions of time may be obtained under the provisions of 37 CFR 1.136(a).

• A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 CFR 1.821(e). If the effective filing date is on or after September 8, 2000, see the final rulemaking notice published in the Federal Register at 65 FR 54604 (September 8, 2000) and 1238 OG 145 (September 19, 2000). Applicant must provide an initial computer readable form (CRF) copy of the "Sequence Listing" and a statement that the content of the sequence listing information recorded in computer readable form is identical to the written (on paper or compact disc) sequence listing and, where applicable, includes no new matter, as required by 37 CFR 1.821(e), 1.821(f), 1.821(g), 1.825(b), or 1.825(d). If applicant desires the sequence listing in the instant application to be identical with that of another application on file in the U.S. Patent and Trademark Office, such request in accordance with 37 CFR 1.821(e) may be submitted in lieu of a new CRF.

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